



SmartPA Criteria Proposal

Drug/Drug Class:	PrEP Fiscal Edit	
First Implementation Date:	April 30, 2020	
Proposed Date:	December 17, 2020	
Prepared for:	MO HealthNet	
Prepared by:	MO HealthNet/Conduent	
Criteria Status:	□Existing Criteria ☑Revision of Existing Criteria □New Criteria	

Executive Summary

Purpose: Ensure appropriate utilization and control of agents used for PrEP

Why Issue Selected:

In his State of the Union Address on February 5, 2019, President Donald J. Trump announced a goal to end the HIV epidemic in the United States within 10 years. The new initiative, named Ending the HIV Epidemic: A Plan for America, seeks to reduce the number of new HIV infections in the United States by 75 percent within five years, and then by at least 90 percent within 10 years, for an estimated 250,000 total HIV infections averted. Pre-exposure prophylaxis (PrEP), a once daily oral regimen antiretroviral therapy, has proven to be highly effective in preventing HIV infection for individuals at high risk, reducing the risk of acquiring HIV by up to 97 percent. The US Preventative Services Task Force (USPSTF) published their recommendations on PrEP in June of 2019, stating persons at high risk of HIV acquisition be offered PrEP with effective antiretroviral therapy. Currently, only an approximate 200,000 of an estimated 1.1 million Americans who are at risk for HIV currently receive PrEP. At this time, 2 agents are FDA approved for use as PrEP - Truvada® (emtricitabine and tenofovir disoproxil fumarate) and Descovy® (emtricitabine and tenofovir alafenamide). Generic versions of Truvada are also now available. MO HealthNet desires to provide PrEP to all qualifying participants, and as such, it is fiscally advantageous for MO HealthNet to establish guidelines for PrEP therapy. This edit will apply only to participants being treated for PrEP and will not affect participants being treated for HIV infection.

Type of Criteria:	☐ Increased risk of ADI	E ☐ Preferred Drug List

Data Sources: ☐ Only Administrative Databases ☐ Databases + Prescriber-Supplied

Setting & Population

- Drug class for review: Agents used for PrEP
- Age range: All appropriate MO HealthNet participants

Approval Criteria

- For treatment of HIV: Please see Antiretrovirals, Treatment Reference Product List
- For use for PrEP:
 - Claim is for brand Truvada OR
 - Clinical consultant review required for therapy with Descovy or generic Truvada

Denial Criteria

Therapy will be denied if all approval criteria are not met
Required Documentation
Laboratory Results: MedWatch Form: Progress Notes: Other:
Disposition of Edit
Denial: Exception code "0683" (Fiscal Edit) Rule Type: CE
Default Approval Period

1 year

References

- TRUVADA® (emtricitabine and tenofovir disoproxil fumarate) tablets, [package insert]. Foster City, CA: Gilead Sciences, Inc.; June 2020.
- DESCOVY® (emtricitabine and tenofovir alafenamide) tablets, [package insert]. Foster City, CA: Gilead Sciences, Inc.; January 2020.
- U.S. Department of Health & Human Services. What is 'Ending the HIV Epidemic: A Plan for America'?. https://www.hiv.gov/federal-response/ending-the-hiv-epidemic/overview. Accessed November 5, 2020.
- Final Recommendation Statement: Prevention of Human Immunodeficiency Virus (HIV) Infection: Preexposure Prophylaxis. U.S. Preventive Services Task Force. June 2019. https://www.uspreventiveservicestaskforce.org/Page/Document/RecommendationStatementFinal/prevention-of-human-immunodeficiency-virus-hiv-infection-pre-exposure-prophylaxis
- IPD Analytics. Rx Insights: Infectious Disease. HIV Update on Treatment Management. November 2019.
- IPD Analytics. Infectious Disease: HIV. Accessed November 5, 2020.